W

11 5

[nucleotide sequence]polynucleotide encoding a peptide of the ONECUT family, a vector Filed comprising [this nucleotide sequence] said polynucleotide, the polypeptide [sequence] encoded by [this nucleotide sequence] said polynucleotide and [/or] a cell line transformed with said vector [and expressing the peptide of the ONECUT family].

- (Amended) The pharmagentical composition as claimed in of claim 1, [characterized in that] wherein the protein of the ONECUT family is an isoform of HNF-6 [in
- (Amended) The pharmaceutical composition [as claimed in of claim 1, its two isoforms]. [characterized in that] wherein the protein of the ONECUT family is OC-2, the amino acid sequence of which is SEQ ID No. 2.
 - (Amended) The [cellular] pharmaceutical composition [as claimed in]of claim 1, [characterized in that]wherein the protein of the ONECUT family is OC-3, the amino acid sequence of which is SEQ ID No 3.
 - (Amended) The pharmaceutical composition as claimed in any one of the preceding claims, characterized in that $\underline{\text{lof Claim 1, wherein}}$ said [nucleotide and polypeptide $sequences \quad are \underline{|polynucleotide|} \quad \underline{a} \quad human \quad \underline{polynucleotide} [\textbf{nucleotide} \quad \textbf{and} \quad \textbf{polypeptide} \\$ sequencel.
 - The pharmaceutical composition [as claimed in any one of the preceding claims, characterized in that of Claim 1, wherein the vector is [chosen]selected from the group consisting of plasmids, viruses, phagemids, lipid vesicles, in particular cationic vesicles, liposomes [or] and a mixture [of these] thereof.
 - (Amended) [The use of the pharmaceutical composition as claimed in any one of the preceding claims, for preparing a medicinal product intended \underline{A} method for the prevention and/or for the treatment of type 1 or type 2 diabetes or of disorders linked to diabetes, for the prevention and/or for the treatment of ancerl, in particular of melanoma, and for the prevention and for the treatment of Waardenburg syndrome, comprising:

administration of the pharmaceutical composition of Claim 1 in an amount effective to prevent or reduce the symptoms of diabetes, cancer, and/or Waardenburg syndrome.

(Amended) [A]The method of [therapeutic treatment of a patient, preferably of a human patient, likely to develop or suffering from diabetes, from a cancer, in particular from a melanoma, or from Waardenburg syndrome, characterized in that |Claim 7, wherein -2-